STRUCTURE UPLOADED L4

=> s 14 sss full FULL SEARCH INITIATED 16:52:33

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SEARCH TIME: 00.00.02

1.5 28 SEA SSS FUL L4

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=> s 15

12 L5 L6

=> d hitstr ibib abs 1-12

ANSWER 1 OF 12 CAPLUS COPYRIGHT 2002 ACS 1.6

IT 263765-27-1P 321883-06-1P 321883-08-3P

RL: PEP (Physical, engineering or chemical process); PYP (Physical process); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (novel lipophilic compds. having affinity with nucleic acids and therapeutical uses thereof)

263765-27-1 CAPLUS RN

CN Phosphonium, [2-[bis(tetradecyloxy)phosphinyl]ethyl]trimethyl-, iodide (9CI) (CA INDEX NAME)

Me- (CH₂)₁₃-O-
$$\frac{O}{\parallel}$$

O- (CH₂)₁₃-Me

• I-

RN 321883-06-1 CAPLUS

CN Phosphonium, [3-[bis(tetradecyloxy)phosphinyl]propyl]trimethyl-, iodide (9CI) (CA INDEX NAME)

Me-
$$(CH_2)_{13}$$
-O- P - $(CH_2)_{3}$ - P +Me₃ O- $(CH_2)_{13}$ - Me

• I-

RN 321883-08-3 CAPLUS

CN Phosphonium, [[bis(tetradecyloxy)phosphinyl]methyl]trimethyl-, iodide (9CI) (CA INDEX NAME)

Me-
$$(CH_2)_{13}$$
-O- P- CH_2 - P+Me₃
O- $(CH_2)_{13}$ - Me

• I-

ACCESSION NUMBER: 200

2001:936089 CAPLUS

DOCUMENT NUMBER:

136:74570

TITLE:

Novel lipophilic compounds having affinity with nucleic acids and therapeutical uses thereof

INVENTOR(S):

Yaouanc, Jean-Jacques; Guenin, Erwann; Clement, Jean-Claude; Herve, Anne-Cecile; Ferec, Claude; Floch,

Virginie; Des Abbayes, Herve

PATENT ASSIGNEE(S):

Fr.

SOURCE:

U.S. Pat. Appl. Publ., 21 pp.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|-----------------------|------|----------|-------------------|--------------|
| - | | | | - |
| US 2001056074 | A1 | 20011227 | US 2001-754814 | 20010104 |
| PRIORITY APPLN. INFO. | : | | US 2000-175342P P | 20000110 |

OTHER SOURCE(S): MARPAT 136:74570

AB The invention consists of a compd. of the general formula R1R2R3A+R4, X-, wherein A, R1, R2, R3, R4 and X are as disclosed in the specification. The invention also relates to the therapeutical uses of this compd., particularly for gene therapy.

L6 ANSWER 2 OF 12 CAPLUS COPYRIGHT 2002 ACS

IT 263765-27-1P 321883-06-1P 321883-07-2P 321883-08-3P 321883-09-4P 321883-10-7P 321883-11-8P 321883-12-9P 321883-13-0P 321883-14-1P 321883-15-2P

RL: ADV (Adverse effect, including toxicity); BPR (Biological process); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses)

(gene delivery by cationic phosphonolipids: effect of structure on transfection)

RN 263765-27-1 CAPLUS

CN Phosphonium, [2-[bis(tetradecyloxy)phosphinyl]ethyl]trimethyl-, iodide (9CI) (CA INDEX NAME)

Me- (CH₂)₁₃-O-
$$\frac{0}{\parallel}$$

P-CH₂-CH₂-P+Me₃
O- (CH₂)₁₃-Me

• I-

RN 321883-06-1 CAPLUS

CN Phosphonium, [3-[bis(tetradecyloxy)phosphinyl]propyl]trimethyl-, iodide (9CI) (CA INDEX NAME)

Me-
$$(CH_2)_{13}$$
-O- P - $(CH_2)_{3}$ - P +Me₃
O- $(CH_2)_{13}$ - Me

• I-

RN 321883-07-2 CAPLUS

CN Phosphonium, [3-[bis(octadecyloxy)phosphinyl]propyl]trimethyl-, iodide (9CI) (CA INDEX NAME)

• I-

RN 321883-08-3 CAPLUS

CN Phosphonium, [[bis(tetradecyloxy)phosphinyl]methyl]trimethyl-, iodide (9CI) (CA INDEX NAME)

Me- (CH₂)₁₃-O-
$$\frac{O}{P}$$
-CH₂- $\frac{P+Me_3}{O}$ -(CH₂)₁₃-Mo

• I-

RN 321883-09-4 CAPLUS

CN Phosphonium, [[bis(tetradecyloxy)phosphinyl]methyl]ethyldimethyl-, iodide (9CI) (CA INDEX NAME)

• I-

RN 321883-10-7 CAPLUS

CN Phosphonium, [[bis(tetradecyloxy)phosphinyl]methyl]dimethylpropyl-, iodide (9CI) (CA INDEX NAME)

• I-

RN 321883-11-8 CAPLUS

CN Phosphonium, [[bis(tetradecyloxy)phosphinyl]methyl]butyldimethyl-, iodide (9CI) (CA INDEX NAME)

• I-

RN 321883-12-9 CAPLUS

CN Phosphonium, [[bis(tetradecyloxy)phosphinyl]methyl]methylbis(1-methylethyl)-, iodide (9CI) (CA INDEX NAME)

• I-

RN 321883-13-0 CAPLUS

CN Phosphonium, [[bis(tetradecyloxy)phosphinyl]methyl]tris(1-methylethyl)-, iodide (9CI) (CA INDEX NAME)

Me-
$$(CH_2)_{13}$$
-O- p - CH_2 - p + $(Pr$ - $i)_3$ | O- $(CH_2)_{13}$ - Me

• I-

RN 321883-14-1 CAPLUS

CN Phosphonium, [[bis(octadecyloxy)phosphinyl]methyl]trimethyl-, iodide (9CI) (CA INDEX NAME)

Me- (CH₂)₁₇-O- P- CH₂- P+Me₃
$$\begin{vmatrix} & & & & & & & & & & \\ & & & & & & & \\ & & & & & & & \\ & & & & & & & \\ & & & & & & & \\ & & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & & \\ & & & \\ & & & & \\ & & & & \\ & & \\ & & & \\ & & \\ & & & \\ & & \\ & & \\ & & & \\ & &$$

I-

RN 321883-15-2 CAPLUS

Phosphonium, [2-[bis(octadecyloxy)phosphinyl]ethyl]trimethyl-, iodide CN (9CI) (CA INDEX NAME)

Me-
$$(CH_2)_{17}$$
-O- P- CH_2 - CH_2 - P+Me3
O- $(CH_2)_{17}$ -Me

ACCESSION NUMBER:

2000:783406 CAPLUS

DOCUMENT NUMBER:

134:136543

TITLE:

Cation Substitution in Cationic Phosphonolipids: A New Concept To Improve Transfection Activity and Decrease

Cellular Toxicity

AUTHOR(S):

Floch, Virginie; Loisel, Severine; Guenin, Erwann; Herve, Anne Cecile; Clement, Jean Claude; Yaouanc, Jean Jacques; des Abbayes, Herve; Ferec, Claude Centre de Biogenetique, CHU ETSBO, Brest, 29275, Fr.

CORPORATE SOURCE: SOURCE:

Journal of Medicinal Chemistry (2000), 43(24),

4617-4628

CODEN: JMCMAR; ISSN: 0022-2623 American Chemical Society

PUBLISHER: DOCUMENT TYPE:

Journal

LANGUAGE:

English

Cationic lipids have been shown to be an interesting alternative to viral vector-mediated gene delivery into in vitro and in vivo model applications. Prior studies have demonstrated that even minor structural modifications of the lipid hydrophobic domain or of the lipid polar domain result in significant changes in gene delivery efficiency. Previously, we developed a novel class of cationic lipids called cationic phosphonolipids and described the ability of these vectors to transfer DNA into different cell lines and in vivo. Up until now, in all new cationic lipids, nitrogen atoms have always carried the cationic or polycationic charge. Recently we have developed a new series of cationic phosphonolipids characterized by a cationic charge carried by a phosphorus or arsenic atom. In a second step, we have also examd. the effects of the linker length between the cation and the hydrophobic domain as regards transfection activity. Transfection activities of this library of new cationic phosphonolipids were studied in vitro in different cell lines (HeLa, CFT1, K562) and in vivo using a luciferase reporter gene. A luminescent assay was carried out to assess luciferase expression. We demonstrated that cation substitution on the polar domain of cationic phosphonolipids (N .fwdarw. P or As) results in significant increase in transfection activity for both in vitro and in vivo assays and decrease of cellular toxicity.

REFERENCE COUNT:

37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 3 OF 12 CAPLUS COPYRIGHT 2002 ACS

263765-27-1P 263765-30-6P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(cationic phosphonolipids contg. quaternary phosphonium and arsonium groups for DNA transfection with good efficiency and low cellular toxicity)

263765-27-1 CAPLUS RN

Phosphonium, [2-[bis(tetradecyloxy)phosphinyl]ethyl]trimethyl-, iodide (9CI) (CA INDEX NAME)

Me-
$$(CH_2)_{13}$$
-O- P- CH_2 - CH_2 - P+Me₃
O- $(CH_2)_{13}$ -Me

) I-

263765-30-6 CAPLUS RN

Phosphonium, [4-[bis(tetradecyloxy)phosphinyl]butyl]trimethyl-, iodide CN (9CI) (CA INDEX NAME)

Me- (CH₂)₁₃-O-P- (CH₂)₄-P+Me₃

$$|$$
O- (CH₂)₁₃-Me

ACCESSION NUMBER:

CORPORATE SOURCE:

2000:132436 CAPLUS

DOCUMENT NUMBER:

132:284071

TITLE:

Cationic phosphonolipids containing quaternary

phosphonium and arsonium groups for DNA transfection

with good efficiency and low cellular toxicity

Guenin, Erwann; Herve, Anne-Cecile; Floch, Virginie; AUTHOR(S):

Loisel, Severine; Yaouanc, Jean-Jacques; Clement,

Jean-Claude; Ferec, Claude; Des Abbayes, Herve

UMR CNRS 6521. Universite de Bretagne Occidentale,

Departement de Chimie, Brest, 29285, Fr.

SOURCE:

Angewandte Chemie, International Edition (2000),

39(3), 629-631

CODEN: ACIEF5; ISSN: 1433-7851

PUBLISHER:

Wiley-VCH Verlag GmbH

DOCUMENT TYPE:

Journal

English

LANGUAGE:

Lipid ammonium, arsonium, and phosphonium compds. such as (C140290)2P(0)(CH2)nA+Me3 I- (A = N, As, and P) were prepd. Phosphonium and arsonium salts are more efficient than ammonium compds. for transfection into Hela cells. Phosphonium and arsonium compds. are less

cytotoxic than the ammonium derivs.

THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 15 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 4 OF 12 CAPLUS COPYRIGHT 2002 ACS

183203-99-8P, Diethyl phosphonomethyltributylphosphonium triflate RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn. of Wittig reagents and their use in prepg. .alpha.,.beta.unsatd. phosphonates)

183203-99-8 CAPLUS RN

Phosphonium, tributyl[(diethoxyphosphinyl)methyl]-, salt with trifluoromethanesulfonic acid (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 183203-98-7 CMF C17 H39 O3 P2

CM 2

37181-39-8 CRN CMF C F3 O3 S

ACCESSION NUMBER:

1999:12341 CAPLUS

DOCUMENT NUMBER:

130:81643

TITLE:

Wittig reagents and method for preparing .alpha.,.beta.-unsaturated phosphonates

Xu, Yibo; Flavin, Michael T.

PATENT ASSIGNEE(S):

Medichem Research, Inc., USA

SOURCE:

INVENTOR(S):

U.S., 8 pp. CODEN: USXXAM

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|-----------------|------|-----------------|-----------------|----------|
| | | | | |
| US 5852198 | А | 19981222 | US 1997-831233 | 19970402 |
| BUED COUDCE/C). | 3.67 | DDATE 120.016/2 | , | |

OTHER SOURCE(S): MARPAT 130:81643

Wittig-type reagents and methods of prepn. and use thereof for prepg. .alpha.,.beta.-unsatd. phosphonate esters from aldehydes and ketones are disclosed. The Wittig-type reagents have the following formulas: (R10)2P(0)CH2P+R3 X- and (R10)2P(0)CH:PR3 wherein X represents triflate, halide, BF4, SbF6, or ClO4; R1 represents alkyl, aryl or arylalkyl; and R represents alkyl, aryl or arylalkyl, provided that R1 and R not represent Ph at the same time. The Wittig reagent di-Et phosphonomethylidenetriphenylphosphorane (1b) was successfully synthesized for the 1st time via its phosphonium triflate salt (4a), by treating di-Et phosphonomethyl triflate with PPh3. The procedure was applied to the synthesis of other new Wittig-type reagents such as phosphoranes and phosphonium salts. The new Wittig reagents thus synthesized were treated with various aldehydes and an activated ketone, affording the corresponding .alpha.,.beta.-unsatd. phosphonates, satd. phosphonates or

phosphoric acids. Triphenylphosphorane 1b and triphenylphosphonium 4a led to both cis and trans isomers with the latter being predominant, while trans isomers were almost exclusively formed when tri-Bu reagents were

REFERENCE COUNT:

32 THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 5 OF 12 CAPLUS COPYRIGHT 2002 ACS L6

IT 205243-74-9P, Tributyl (2-dimethoxyphosphinyl-2-

(methylene)ethyl)phosphonium chloride

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn. and thermolytic conversion to betaine)

205243-74-9 CAPLUS RN

Phosphonium, tributyl[2-(dimethoxyphosphinyl)-2-propenyl]-, chloride (9CI) CN(CA INDEX NAME)

● cl-

ACCESSION NUMBER:

1998:224959 CAPLUS

DOCUMENT NUMBER:

128:257508

TITLE:

Betaine formation from dimethyl 3-phosphoniopropen-2-

ylphosphonate chlorides

AUTHOR(S):

Gurevich, I. E.; Tebby, J.; Dogadina, A. V.; Ionin, B.

CORPORATE SOURCE:

Staffordshire University, Stoke-on-Trent, UK

SOURCE:

Russian Journal of General Chemistry (Translation of

Zhurnal Obshchei Khimii) (1997), 67(2), 324-325

CODEN: RJGCEK; ISSN: 1070-3632

PUBLISHER:

MAIK Nauka/Interperiodica Publishing

DOCUMENT TYPE:

Journal

LANGUAGE:

English

(MeO)2P(O)C(CH2Cl):CH2 reacts with PBuPh2 and PBu3 to give AΒ

[(MeO)2P(O)C(:CH2)CH2PR12R2]C1 (R1/R2 = Ph/Bu, Bu/Bu), which form betaines (MeO)P(O)(O-)C(:CH2)CH2PR12R2+ on heating without isomerization in CHCl3 or THF.

L6 ANSWER 6 OF 12 CAPLUS COPYRIGHT 2002 ACS

IT 183203-99-8P

> RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn., deprotonation, or reaction with aldehydes or ketones)

RN 183203-99-8 CAPLUS

Phosphonium, tributyl[(diethoxyphosphinyl)methyl]-, salt with CN trifluoromethanesulfonic acid (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 183203-98-7 CMF C17 H39 O3 P2

CM 2

CRN 37181-39-8 CMF C F3 O3 S

F-C-so3-

ACCESSION NUMBER:

1997:679095 CAPLUS

DOCUMENT NUMBER:

127:319076

TITLE:

Wittig reagents and method for preparing .alpha.,.beta.-unsaturated phosphonates Xu, Yibo; Flavin, Michael T.; Xu, Ze-Qi

INVENTOR(S):
PATENT ASSIGNEE(S):

Medichem Research, Inc., USA; Xu, Yibo; Flavin,

Michael T.; Xu, Ze-Qi PCT Int. Appl., 25 pp.

SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

```
PATENT NO.
                   KIND
                           DATE
                                          APPLICATION NO. DATE
                    A1
                           19971009
                                        WO 1997-US5474 19970402
        W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
            DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC,
            LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
            RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN,
            AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB,
            GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN,
            ML, MR, NE, SN, TD, TG
    AU 9724348
                      A1
                         19971022
                                          AU 1997-24348
                                                           19970402
                                       US 1996-14764P P 19960403
PRIORITY APPLN. INFO.:
                                                       W 19970402
                                       WO 1997-US5474
```

OTHER SOURCE(S):

CASREACT 127:319076; MARPAT 127:319076

AB The prepn. of Wittig-type reagents, (R10)2P(0)CH2P+(R2)3X- and (R10)2P(0)CH:P(R2)3 (X = triflate, halide, BF4, SbF6, ClO4; R1 = alkyl, aryl, arylalkyl; R2 = alkyl, aryl, arylalkyl, provided that R1 and R2 do not represent Ph at the same time), and use thereof, starting from aldehydes and ketones are disclosed. The Wittig reagent di-Et phosphonomethylidenetriphenylphosphorane has been successfully synthesized for the first time via its phosphonium triflate salt [{(EtO)2P(O)CH2PPh3}+(OTf)-], by treating di-Et phosphonomethyltriflate with PPh3 according to the disclosed method. The procedure has been applied to the synthesis of other new Wittig-type reagents such as phosphoranes and phosphonium salts. The new Wittig reagents thus synthesized were treated with various aldehydes and an activated ketone,

affording the corresponding .alpha.,.beta.-unsatd. phosphonates, satd. phosphonates or phosphoric acids. Triphenylphosphorane and triphenylphosphonium led to both cis and trans isomers with the latter being predominant, while trans isomers were almost exclusively formed when tri-Bu reagents were used.

L6 ANSWER 7 OF 12 CAPLUS COPYRIGHT 2002 ACS

IT 183203-99-8P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn. of new Wittig reagents and application to synthesis of unsatd. phosphonates)

RN 183203-99-8 CAPLUS

CN Phosphonium, tributyl[(diethoxyphosphinyl)methyl]-, salt with trifluoromethanesulfonic acid (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 183203-98-7 CMF C17 H39 O3 P2

CM 2

CRN 37181-39-8 CMF C F3 O3 S

ACCESSION NUMBER: 1996:618940 CAPLUS

DOCUMENT NUMBER: 125:328879

TITLE: Preparation of New Wittig Reagents and Their

Application to the Synthesis of .alpha.,.beta.-

Unsaturated Phosphonates

AUTHOR(S): Xu, Yibo; Flavin, Michael T.; Xu, Ze-Qi

CORPORATE SOURCE: MediChem Research Inc., Lemont, IL, 60439, USA

SOURCE: Journal of Organic Chemistry (1996), 61(22), 7697-7701

CODEN: JOCEAH; ISSN: 0022-3263

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 125:328879

AB The Wittig reagent [(diethoxyphosphinyl)methylidene]triphenylphosphorane (1b) has been successfully synthesized for the first time via its phosphonium triflate salt (EtO)2P(O)CH2P+Ph3 OTf- (4a), by treating (diethoxyphosphinyl)methyl triflate with triphenylphosphine. The procedure has been applied to the synthesis of other phosphoranes and phosphonium salts. The new Wittig reagents thus synthesized were treated

with various aldehydes and an activated ketone, affording the corresponding .alpha.,.beta.-unsatd. phosphonates. Triphenylphosphorane 1b and triphenylphosphonium 4a led to both cis and trans isomers with the latter being predominant, while trans isomers were almost exclusively formed when tri-Bu reagents were used.

L6 ANSWER 8 OF 12 CAPLUS COPYRIGHT 2002 ACS

IT 117740-93-9P 117740-94-0P 117740-95-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (prepn. and thermal decompn. of)

RN 117740-93-9 CAPLUS

CN Phosphonium, [3-chloro-2-(diethoxyphosphinyl)-2-propenyl]trimethyl-, chloride (9CI) (CA INDEX NAME)

● cl-

RN 117740-94-0 CAPLUS

CN Phosphonium, [3-chloro-2-(diethoxyphosphinyl)-2-propenyl]tripropyl-, chloride (9CI) (CA INDEX NAME)

● c1-

RN 117740-95-1 CAPLUS

CN Phosphonium, tributyl[3-chloro-2-(diethoxyphosphinyl)-2-propenyl]-, chloride (9CI) (CA INDEX NAME)

● c1-

ACCESSION NUMBER:

1988:631163 CAPLUS

DOCUMENT NUMBER: 109:231163

TITLE: Quaternary phosphonium salts in the vinyl phosphate

series

AUTHOR(S):

Gololobov, Yu. G.; Oganesyan, A. S.; Petrovskii, P. V.

CORPORATE SOURCE:

Inst. Elementoorg. Soedin. im. Nesmeyanova, Moscow,

USSR

SOURCE:

Zh. Obshch. Khim. (1988), 58(1), 225-6

CODEN: ZOKHA4; ISSN: 0044-460X

DOCUMENT TYPE:

Journal

LANGUAGE:

. Russian

OTHER SOURCE(S):

CASREACT 109:231163

Treating (EtO)2P(O)OC(:CHCl)CH2Cl with R3P (R = Me, Pr, Bu) at 0.degree. for 2 days gave (EtO)2P(O)OC(:CHCl)CH2PR3+ Cl- (same R) quant., which gave betaines R3P+CH2C(:CHCl)OP(O)(OEt)O- on heating.

ANSWER 9 OF 12 CAPLUS COPYRIGHT 2002 ACS L6

IT88410-11-1P

RL: SPN (Synthetic preparation); PREP (Preparation)

(prepn. of)

88410-11-1 CAPLUS RN

CN Phosphonium, tributyl[(diethoxyphosphinyl)difluoromethyl]-, bromide (9CI)

● Br-

ACCESSION NUMBER:

1984:51710 CAPLUS

DOCUMENT NUMBER:

100:51710

TITLE:

Preparation and synthetic utility of fluorinated

phosphonium salts, bisphosphonium salts and

phosphoranium salts

AUTHOR(S):

Burton, Donald J.

CORPORATE SOURCE:

Dep. Chem., Univ. Iowa, Iowa City, IA, 52242, USA

SOURCE: J. Fluorine Chem. (1983), 23(4), 339-57

CODEN: JFLCAR; ISSN: 0022-1139

DOCUMENT TYPE:

Journal

LANGUAGE:

French

The reaction of PPh3,P(NMe2)3, and PBu3 with CF2Br2, CF2BrI, CFBr3, and CFC13 gave rapid and high yield synthesis of various types of fluorinated phosphonium salts, bisphosphonium salts and phosphoranium salts. These salts are useful precursors to fluorine-contg. ylides, carbenes and methide ions, and were used to prep. 1,1-dihaloalkenes from ketones or aldehydes. The prepn., mechanism of formation, and the synthetic utility of these novel reagents were described.

ANSWER 10 OF 12 CAPLUS COPYRIGHT 2002 ACS L6

79443-90-6P 79443-91-7P IT

> RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of)

RN 79443-90-6 CAPLUS

Phosphonium, tributyl[2-chloro-4-(diethoxyphosphinyl)-2-butenyl]-, bromide (9CI) (CA INDEX NAME)

● Br-

RN 79443-91-7 CAPLUS

CN Phosphonium, tributyl[2-chloro-4-(diethoxyphosphinyl)-1-butenyl]-, bromide (9CI) (CA INDEX NAME)

● Br=

ACCESSION NUMBER:

1981:569300 CAPLUS

DOCUMENT NUMBER:

95:169300

TITLE:

Reactions of phosphonium salts and phosphonate

obtained from 1,4-dibromo-2-chloro-2-butene

AUTHOR(S):

Lulukyan, R. K.; Ovakimyan, M. Zh.; Panosyan, G. A.;

Indzhikyan, M. G.

CORPORATE SOURCE:

SOURCE:

Inst. Org. Khim., Yerevan, USSR
Arm. Khim. Zh. (1981), 34(6), 474-9

CODEN: AYKZAN; ISSN: 0515-9628

DOCUMENT TYPE:

Journal

LANGUAGE:

Russian

AB Mono- and diphosphonium salts with a Cl atom at the .gamma.-position [e.g., R3P+CH2CH:CClCH2R1 Br- (R = Ph, Bu; R1 = Br, PR3+ Br-)] and (EtO)2P(O)CH2CH:CClCH2Br were prepd. by appropriate reactions of BrCH2CH:CClCH2Br, and their nucleophilic substitutions with phosphines and amines were investigated.

L6 ANSWER 11 OF 12 CAPLUS COPYRIGHT 2002 ACS

IT 54580-38-0P 54580-42-6P 54580-59-5P

RN 54580-38-0 CAPLUS

CN Phosphonium, [2-(butoxymethoxyphosphinyl)ethyl]trimethyl-, bromide (9CI) (CA INDEX NAME)

$$\begin{array}{c} \text{O} \\ \parallel \\ \text{n-BuO-} \begin{array}{c} \text{P-} \text{CH}_2\text{--} \text{CH}_2\text{--} \text{P+Me}_3 \\ \parallel \\ \text{OMe} \end{array}$$

● Br-

RN 54580-42-6 CAPLUS

CN Phosphonium, [4-(butoxymethoxyphosphinyl)-4-oxobutyl]trihexadecyl-, bromide (9CI) (CA INDEX NAME)

● Br-

RN 54580-59-5 CAPLUS

CN Phosphonium, butyl[3-(diethoxyphosphinyl)-3-ethyl-4-oxopentyl]ethylhexyl-, chloride (9CI) (CA INDEX NAME)

• C1-

ACCESSION NUMBER:

1975:31392 CAPLUS

DOCUMENT NUMBER:

82:31392

TITLE:

Organophosphonium salts

INVENTOR(S):

Grayson, Martin; Keough, Patricia T.

PATENT ASSIGNEE(S):

American Cyanamid Co.

SOURCE:

U.S., 9 pp. Division of U.S. 3,689,601 (CA

77;152346v). CODEN: USXXAM

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

. 2

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------|------|----------|-----------------|----------|
| | | | | |
| US 3836587 | Α | 19740917 | US 1972-255770 | 19720522 |

| US 3689 | 601 | Α | 19720905 |
|--------------|-----------|---|----------|
| PRIORITY APP | LN. INFO. | : | |

US 1969-871628 19691117 US 1963-292123 19630701 US 1967-674107 19671010 US 1969-871628 19691117

AB The reaction of R3P with XCH2CH2OH gave .apprx.20 R3P+CH2CH2OH X- [R = Me, Et, Bu, Ph, cyclohexyl, p-tolyl, Me(CH2)15, etc.; X = Cl, Br, iodo], which were esterified and(or) dehydrated to give CH2:CHP+R3 X-. These vinyl phosphonium salts condensed with compds. such as Me2CO, CH2CO2Me, cyclohexanone, indene, MeNO2, Ph2PHO, etc., to give phosphinoethyl derivs.

L6 ANSWER 12 OF 12 CAPLUS COPYRIGHT 2002 ACS

IT 23685-59-8P 23685-60-1P 23756-92-5P

RN 23685-59-8 CAPLUS

CN Phosphonium, diheptyl(hydroxymethyl)(phosphonomethyl)-, chloride, dioctyl ester (8Cİ) (CA INDEX NAME)

● Cl-

RN 23685-60-1 CAPLUS

CN Phosphonium, diheptyl(hydroxymethyl)(2-phosphonoethyl)-, chloride, dioctyl ester (8CI) (CA INDEX NAME)

● cl-

RN 23756-92-5 CAPLUS

CN Phosphonium, dibutyl(hydroxymethyl)(phosphonomethyl)-, chloride, dihexyl ester (8CI) (CA INDEX NAME)

ACCESSION NUMBER:

1969:470693 CAPLUS

DOCUMENT NUMBER:

TITLE:

Preparation of dialkylethoxycarbonylmethylphosphines

and phosphine oxides

AUTHOR(S): CORPORATE SOURCE: Petrov, K. A.; Parshina, V. A.; Petrova, G. M.

SOURCE:

Zh. Obshch. Khim. (1969), 39(6), 1247-51

CODEN: ZOKHA4

DOCUMENT TYPE:

Journal

71:70693

LANGUAGE:

Russian

Heating an equimolar mixt. of R2PCH2OH and XCH2CO2Et or its analogs, where X = Cl or Br, in an inert atm. at up to 130.degree., finally in vacuo, several hrs. gave the following R2R1PCH2OH+X- (R, R1, X, and n20D given): Bu, Ch2CO2Et, Cl, 1.5020; Bu, CH2CO2Et, Br (I), 1.5010; Bu, CH2P(O)Bu2, Cl, 1.4850; Bu, CH2P(O)(OC6H13)2, Cl, 1.4696; C7H15, CH2CO2Et, Br, 1.4964; C7H15, CH2P(O)(OC8H17)2, Cl, 1.4771; and C7H15, CH2CH2P(O)(OC8H17)2, Cl, 1.4855. The products were purified with activated C. Treating I 4 hrs. at 80.degree. with Et3N gave 26% Bu2PCH2CO2Et, b0.02-0.03 150-60.degree. (bath), n21.5D 1.4625, insol. in Et20. Similarly was prepd. the diheptyl analog, n18.5D 1.4650; similar reaction with N2H4 gave (Bu2PCH2CONH)2, an oil. Oxidn. of the above phosphines with 10% H2O2 gave the oxides: Bu2P(O)CH2CO2Et, b0.02-0.03 105-7.degree.; diheptyl analog, b0.02-0.03 112.degree., n21.5D 1.4620. ClCH2POCl2 (41.9 g.) added to 78.5 g. C8H170H and 40 ml. pyridine in C6H6 gave after 3 hrs. heating 50.5%ClCH2P(O)(OC8H17)2, b2 185-8.degree., n17D 1.4511. Similarly was prepd. the 2-chloroethylphosphonate analog, b1 182-5.degree., n22D 1.4480. Ir spectral data were given.



| L Number | Hits | Search Text | DB | Time stamp |
|----------|------|---|-----------|------------------|
| 1 | 0 | cationic and phosphonium and phosphonolipid | USPAT; | 2002/08/20 17:45 |
| j j | | | US-PGPUB; | |
| | | | EPO; JPO; | |
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| 2 | 1 | cationic and phosphonolipid | USPAT; | 2002/08/20 17:45 |
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| 3 | 0 | phosphonium and phosphonolipid | USPAT; | 2002/08/20 17:50 |
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| 4 | 6715 | phosphonium and cationic | USPAT; | 2002/08/20 17:46 |
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| 7 | 115 | (((phosphonium and cationic) and lipid) and phosphon\$) | USPAT; | 2002/08/20 17:46 |
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| 8 | 49 | ((((phosphonium and cationic) and lipid) and phosphon\$) | USPAT; | 2002/08/20 17:47 |
| | | and dna) and quatern\$ | US-PGPUB; | |
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| 9 | 7 | dna and phosphonolipid | USPAT; | 2002/08/20 17:50 |
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| - | 0 | yaouanac-\$.in. | USPAT; | 2002/08/20 17:42 |
| | | | US-PGPUB; | |
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| - | 5 | yaouanc-\$.in. | USPAT; | 2002/08/20 13:02 |
| | | | US-PGPUB; | |
| | | | EPO; JPO; | |
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| - | 3 | Petrov-\$.in. and phosphine | USPAT; | 2002/08/20 17:09 |
| | | | US-PGPUB; | |
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| - | 1 | 1984-224528.NRAN. | DERWENT | 2002/08/20 16:33 |
| - | 9 | "5674908" | USPAT; | 2002/08/20 17:34 |
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(FILE 'HOME' ENTERED AT 16:36:16 ON 20 AUG 2002)

FILE 'CAPLUS' ENTERED AT 16:36:29 ON 20 AUG 2002 L1 STRUCTURE UPLOADED S L1

FILE 'REGISTRY' ENTERED AT 16:39:57 ON 20 AUG 2002 L2 11 S L1 SSS FULL

FILE 'CAPLUS' ENTERED AT 16:39:58 ON 20 AUG 2002 L3 6 S L2 SSS FULL

FILE 'STNGUIDE' ENTERED AT 16:41:22 ON 20 AUG 2002

FILE 'REGISTRY' ENTERED AT 16:52:22 ON 20 AUG 2002 STRUCTURE UPLOADED

L5 28 S L4 SSS FULL

FILE 'CAPLUS' ENTERED AT 16:52:59 ON 20 AUG 2002 L6 12 S L5

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L1 STR

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L4 STR

Structure attributes must be viewed using STN Express query preparation.